

## Adult Immunization Update 2014

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Immunization Services Division



## Disclosures

- ❑ Dr. Kroger is a federal government employee with no financial interest or conflict with the manufacturer of any product named in the following presentations
- ❑ The off-label use of IIV, MMR, HPV and PCV13 vaccines will be discussed
- ❑ Vaccines not currently licensed by the FDA will not be discussed

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## Take Out Your Cell Phone



- ❑ You will have the opportunity to text answers to questions!
- ❑ Please note: message and data rates may apply

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## What's New in Adult Immunization

- ❑ Disease burden
- ❑ Schedules
- ❑ Adult vaccination coverage rates
- ❑ Pneumococcal vaccines
- ❑ Influenza vaccines
- ❑ Tdap and pregnancy
- ❑ Zoster vaccine
- ❑ Hepatitis B vaccine and HCP
- ❑ Immunization resources

## Burden of Disease Among U.S. Adults for Vaccine-preventable Diseases

- ❑ **Influenza disease burden varies year to year<sup>1</sup>**
  - Millions of cases and average of 226,000 hospitalizations annually with more than 75% among adults
  - 3,000-49,000 deaths annually, more than 90% among adults
- ❑ **Invasive pneumococcal disease (IPD)<sup>2</sup>** - 13,000 invasive pneumococcal disease cases among adults 65 and older in 2013
- ❑ **Pertussis<sup>3</sup>** - 41,880 total reported cases 2012 (~9,000 among adults)
- ❑ **Hepatitis B<sup>4</sup>** - 3,350 acute cases reported 2010
- ❑ **Zoster<sup>5</sup>** - ~1 million cases annually

<sup>1</sup>CDC. Estimates of deaths associated with seasonal influenza United States, 1976-2007. *MMWR* 2010; 59(33):1057-1062.

<sup>2</sup>CDC. Active Bacterial Core Surveillance. [www.cdc.gov/ncez/reports\\_findings/surveillance/special/13.pdf](http://www.cdc.gov/ncez/reports_findings/surveillance/special/13.pdf)

<sup>3</sup>CDC. Notifiable Diseases and Morbidity Tables. *MMWR* 2013; 61(18):450-451.

<sup>4</sup>CDC. Viral Hepatitis Surveillance United States, 2010. National Center for HIV/AIDS, Viral Hepatitis, STD & TB Prevention/Division of Viral Hepatitis.

<sup>5</sup>CDC. Prevention of Herpes Zoster. *MMWR* 2008; 57(6):1-10.

## HPV and HPV Vaccine-preventable Cancers in Women and Men, 2005-2009

Anatomic Area	Average annual number of cases <sup>a</sup>	Attributable to HPV+ <sup>b</sup>	Attributable to HPV 16/18+ <sup>c</sup>
Cervix	11,279	10,150	7,470
Vagina	694	520	300
Vulva	3,039	2,100	1,480
Anus and rectum	3,064	2,810	2,450
Oropharynx	2,317	1,670	1,420
<b>Total (Women)</b>	<b>21,342</b>	<b>17,250</b>	<b>13,200</b>
Penis	1,003	830	400
Anus and Rectum	1,687	1,540	1,340
Oropharynx	9,312	6,700	5,730
<b>Total (Men)</b>	<b>13,446</b>	<b>8,880</b>	<b>7,550</b>

Jemal A et al. *J Natl Cancer Inst* 2013;105:491

Data presented by Saraiya M, 28<sup>th</sup> International Papillomavirus Conference 2012, Puerto Rico











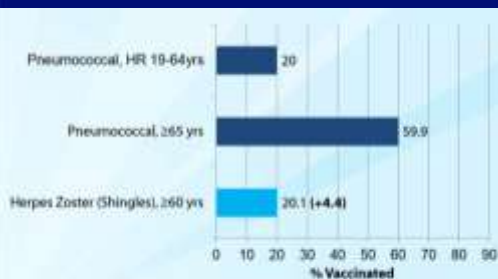


### Influenza Vaccination Coverage among U.S. Adults: 2011-12, 2012-13, and 2013-14 Seasons

Group	2011-12 (%)	2012-13 (%)	2013-14 (%) <sup>a</sup>	Difference (%)
Persons ≥ 18 yrs	38.8	41.5	42.4	3.6
Persons 18-49 yrs, all	28.6	31.1	32.3	3.7
Persons 18-49 yrs, high risk	36.8	39.8	38.7	1.9
Persons 50-64 yrs	42.7	45.1	45.3	2.6
Persons ≥ 65 yrs	64.9	66.2	65.0	0.1

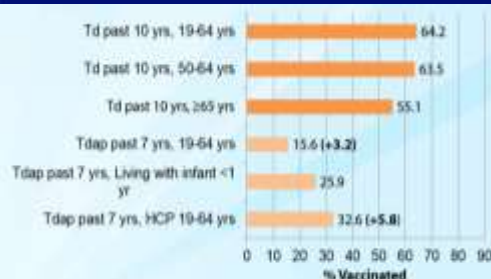
<sup>a</sup> Estimates of the percentage of people vaccinated are based on interviews conducted beginning September (2011-12) or October (2012-13) through June 2013 and reported vaccinations from July 2011 through May 2013. For California, 2013-14 interview data were only available for September-December 2013 and this estimate for persons ≥ 18 years only reflect vaccinations during only November 2013. For Mississippi, sample size was insufficient for persons ≥ 18 years only reflect vaccinations during only November 2013. For Mississippi, sample size was insufficient for persons ≥ 18 years only reflect vaccinations during only November 2013. For Mississippi, sample size was insufficient for persons ≥ 18 years only reflect vaccinations during only November 2013. For Mississippi, sample size was insufficient for persons ≥ 18 years only reflect vaccinations during only November 2013.

### Adult Vaccination Coverage, Selected Vaccines by Age and High-risk Status, United States



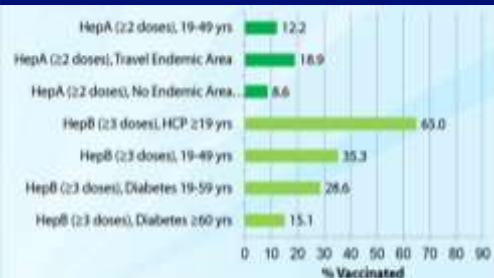
HP2020 Targets: 90% PPV ≥65 yrs, 60% PPV HR 19-64 yrs, 30% Shingles  
Data Source: 2012 NHIS

### Adult Vaccination Coverage, Tetanus-containing by Age and High-risk Status, United States



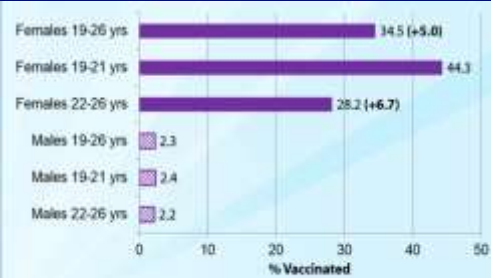
Data Source: 2012 NHIS

### Adult Vaccination Coverage, Hepatitis A and B Vaccines by Age and High-risk Status, United States



HP2020 Target 90% HepB Healthcare Personnel (HCP)  
Data Source: 2012 NHIS

### HPV Vaccination Coverage (1 or more dose ever) Adults 19-26 Years by Sex, United States



Data Source: 2012 NHIS





## UPDATED ACIP RECOMMENDATIONS FOR ADULTS

<http://www.cdc.gov/vaccines/acip/index.html>



## PNEUMOCOCCAL VACCINES

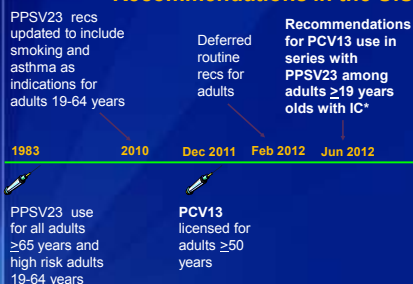
[www.cdc.gov/vaccines/pubs/ACIP-list.htm#pcv](http://www.cdc.gov/vaccines/pubs/ACIP-list.htm#pcv)

### Pneumococcal Vaccines

Characteristic	23-valent Pneumococcal Polysaccharide Vaccine (PPV23)	13-Valent Pneumococcal Conjugate Vaccines (PCV13)
Components	Purified polysaccharide	Purified polysaccharide covalently bound to carrier protein
Ages	2 years and older	6 weeks and older *
Number of serotypes	23	13
Effect against bacteremia	Substantial	Substantial
Effect against carriage	None	Substantial
Effect against non-bacteremic pneumonia	No consensus	Moderate
Route	Subcut or IM injection	IM Injection

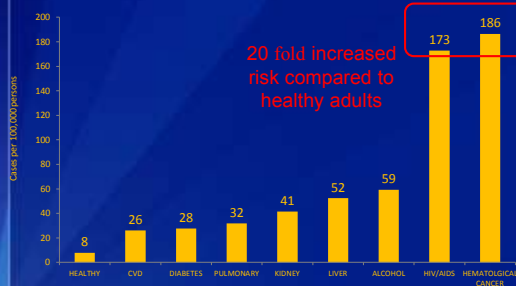
\*ACIP Off-label recommendation

### Timeline for Adult Pneumococcal Vaccine Recommendations in the U.S.



\*IC = with immunocompromising conditions, functional or anatomic asplenia, CSF leaks, or cochlear implants

### Incidence of IPD in Adults Aged 18-64 Years with Selected Underlying Conditions, US, 2009

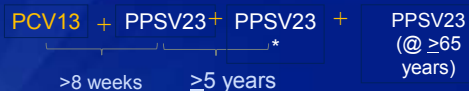


Kyaw, JID 2005;192:377-86 & CDC Unpublished

### Prevention of Pneumococcal Disease Among Adults with Immunocompromising Conditions

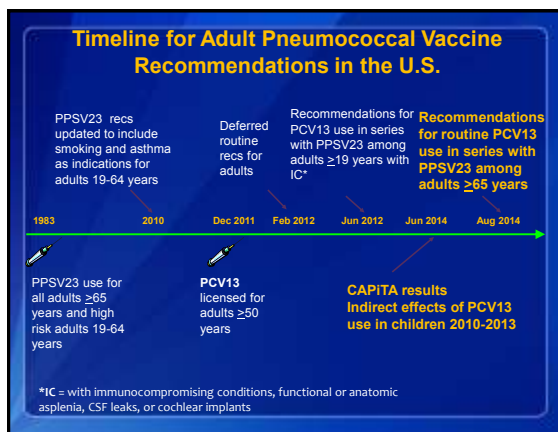
#### Recommendation for PPSV23-naïve adults

Adults 19 years of age or older with immunocompromising conditions, functional or anatomic asplenia, CSF leaks, or cochlear implants, and who have not previously received PCV13 or PPSV23 should receive a dose of PCV13 first followed by currently recommended doses of PPSV23



\*Second PPSV dose before age 65 years NOT recommended for adults with CSF leaks or those with cochlear implants





### Summary of Evidence Supporting PCV13 Use among Adults ≥65 Years of Age

- Results of a large trial demonstrated PCV13 prevents IPD and non-bacteremic pneumonia<sup>1</sup>
  - 75% reduction in vaccine type IPD
  - 45% reduction in vaccine type non-bacteremic pneumonia
- Safety demonstrated in clinical trials<sup>1,2</sup>
- Vaccine preventable disease burden remaining among adults ≥65 years
  - Estimated 2,600 PCV13 type IPD cases in 2013<sup>3</sup>
  - Over 50,000 PCV13-type inpatient CAP<sup>4</sup>

<sup>1</sup>CAPITA, June 2014 ACIP  
<sup>2</sup>Phase III trials, Pfizer, ACIP 2011, 2012  
<sup>3</sup>Active Bacterial Core Surveillance, 2013  
<sup>4</sup>CAP = Community acquired pneumonia

### Summary of Evidence Supporting PCV13 Use in Series with PPSV23 Among Adults ≥65 Years of Age

- In the short-term, PCV13 likely provides adequate coverage of disease causing serotypes
  - 20-25% IPD due to PCV13 types<sup>1</sup>
  - ~10% of all CAP due to PCV13 types<sup>2</sup>
- Broader protection is expected to be provided through use of both PCV13 and PPSV23 in series
  - 38% of IPD due to serotypes unique to PPSV23<sup>1</sup>
- Immune response improved when PCV13 given before PPSV23 in series
- Optimal interval determined based on immune response, safety, the risk window for protection against PPSV-only types, practical considerations

<sup>1</sup>Active Bacterial Core Surveillance, 2013  
<sup>2</sup>Estimate based on studies using serotype-specific urine antigen test, Pfizer.  
 CAP = Community acquired pneumonia

### ACIP Recommendations for PPSV23 for Adults 19 Years and Older 2010

- Immunocompetent w/underlying condition
  - PPSV23 + PPSV23 (@ ≥65 years) ≥5 years
- Asplenic (sickle cell, hemoglobinopathy), Immunocompromised
  - PPSV23 + PPSV23 + PPSV23 (@ ≥65 years) ≥5 years, ≥5 years
- 65 years or older with no high-risk condition
  - PPSV23

### ACIP Recommendations for PCV13 and PPSV23 for Adults 19 Years and Older 2012

- Immunocompromised, Asplenic (sickle cell, hemoglobinopathy), CSF leaks, Cochlear Implants who are **Pneumococcal-naïve**
  - PCV13 + PPSV23 + PPSV23\* + PPSV23 (@ ≥65 years)
  - ≥8 weeks, ≥5 years, ≥5 years

\*Second PPSV23 dose before age 65 years NOT recommended for adults with CSF leaks or those with cochlear implants

\*ACIP off-label recommendation for PCV13 for adults 19 through 49 years of age

### ACIP Recommendations for PCV13 and PPSV23 for Adults 19 Years and Older 2012

- Immunocompromised, Asplenic (sickle cell, hemoglobinopathy), CSF leaks, Cochlear Implants who have previously received PPSV23
  - PPSV23 + PCV13 + PPSV23\* + PPSV23 (@ ≥65 years)
  - ≥1 year, ≥8 weeks, ≥5 years
  - ≥5 years

\*Second PPSV23 dose before age 65 years NOT recommended for adults with CSF leaks or those with cochlear implants

\*ACIP off-label recommendation for PCV13 for adults 19 through 49 years of age



## ACIP Recommendations for PCV13 and PPSV23 for Adults 65 Years and Older 2014

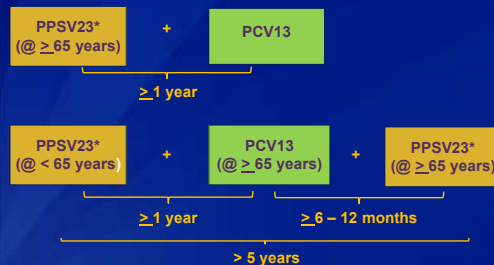
- Pneumococcal-naïve or unknown vaccination history



- If PPSV23 cannot be given during 6-12 month window after PCV13, a dose of PPSV23 should be given during the next visit
- Minimum interval = 8 weeks

## ACIP Recommendations for PCV13 and PPSV23 for Adults 65 Years and Older 2014

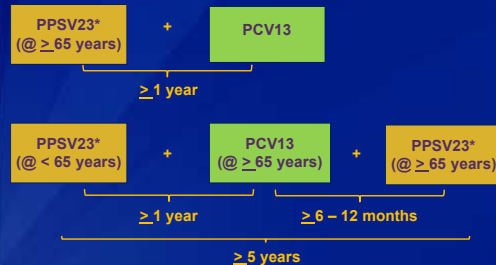
- Previously received one or more doses of PPSV23



\*Doses already administered

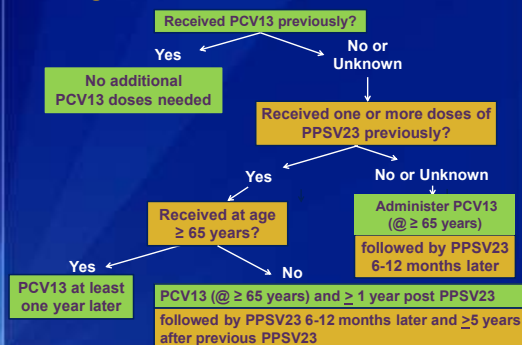
## ACIP Recommendations for PCV13 and PPSV23 for Adults 65 Years and Older 2014

- Previously received one or more doses of PPSV23



\*Doses already administered

## Categories of Adults 65 Years or Older



## Administering PCV13 and PPSV23 Vaccines General Rules

- PCV13 and PPSV23 should not be administered during the same clinic visit
  - Either vaccine may be administered with other vaccines
- Administer PCV13 before PPSV23 whenever possible

## Michigan Provider Resources



“Quick Looks”

[www.michigan.gov/documents/mdchPPSV23\\_PCV13\\_Adults\\_101512\\_Final\\_401240\\_7.pdf](http://www.michigan.gov/documents/mdchPPSV23_PCV13_Adults_101512_Final_401240_7.pdf)



## What Do You Think?



A 66-year-old patient has had laboratory confirmed pneumococcal pneumonia. She has never been vaccinated with pneumococcal vaccine. She should receive:

- A. PPSV23 only - Text IZ1A to 22333
- B. PCV13 only - Text IZ1B to 22333
- C. PCV13 now followed by PPSV23 - Text IZ1C to 22333
- D. PPSV23 now followed by PCV13 – Text IZ1D to 22333

Note: Message and Data Rates May Apply [Poll Results](#)



## PERTUSSIS VACCINATION OF PREGNANT WOMEN

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/tdap-td.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/tdap-td.html)

## ACIP Conclusions: Tdap and Pregnancy

- ❑ Maternal antibodies from women immunized before pregnancy waned quickly (Healy 2012)
  - Concentration of maternal antibodies unlikely high enough to provide passive protection to infants
- ❑ A single dose of Tdap during one pregnancy is insufficient to provide protection for subsequent pregnancies

## Bottom Line

An infant's first dose of pertussis vaccine is the one you administer to his/her mom!

## ACIP Recommendations: Tdap and Pregnancy

- ❑ Administer Tdap to pregnant women during each pregnancy, regardless of previous Tdap vaccination history
- ❑ Ideally vaccinate between 27 through 36 weeks gestation although Tdap may be given at any time during pregnancy
  - 27 through 36 weeks gestation is optimal timing to maximize the maternal antibody response AND passive antibody transfer to the infant

\*ACIP off-label recommendation, MMWR, Vol. 62 No. 7, Feb, 22, 2013

## Special Situations and Pregnant Women

- ❑ Unknown or incomplete tetanus vaccination: should complete the 3-dose primary series
  - Recommended schedule is 0, 4 weeks, and 6 through 12 months
  - Tdap should replace 1 dose of Td, preferably between 27- 36 weeks gestation
- ❑ Wound care: previously unvaccinated pregnant woman should be given Tdap if Td is indicated for wound management

\*ACIP off-label recommendation: MMWR, Vol. 62 No. 7, February 22, 2013 and MMWR 2011;60 (No.41): 11424-1426



## Tdap and Postpartum Women

- ❑ Postpartum women **not previously vaccinated** should be vaccinated immediately
  - Including women who are breastfeeding
- ❑ Do not administer Tdap to postpartum women who have already been vaccinated with Tdap
  - Regardless of the length of time since Tdap vaccination

## Resources for Providers and Pregnant Women



<http://www.cdc.gov/vaccines/hcp/patient-ed/adults-for-patients/pregnant-women.html>

## What Do You Think?



Your patient just delivered her first child. She was previously vaccinated with Tdap as an adolescent. She did not receive Tdap during this pregnancy. Should you administer Tdap vaccine prior to discharge?

A. Yes – Text IZ2Y to 22333

B. No – Text IZ2N to 22333

Note: Message and Data Rates May Apply [Poll Results](#)

## Seasonal Influenza

- ❑ Updated ACIP recommendations were published August 2014

- ❑ Includes:
  - Vaccine composition
  - Vaccination schedules for those
    - 6 months thru 8 years
    - 9 years and older
  - Guidance on product preference



[www.cdc.gov/mmwr/pdf/wk/mm6332.pdf](http://www.cdc.gov/mmwr/pdf/wk/mm6332.pdf)

## the benefits of flu vaccination

The estimated number of influenza-associated illnesses prevented by flu vaccination during the 2012-2013 season:

**6.6 million**



or the population of the state of Arizona

The estimated number of flu-associated medical visits prevented by vaccination during the 2012-2013 season:

**3.2 million**



or the passengers of 1,067 mega-cruise ships

The estimated number of flu hospitalizations prevented during the 2012-2013 season:

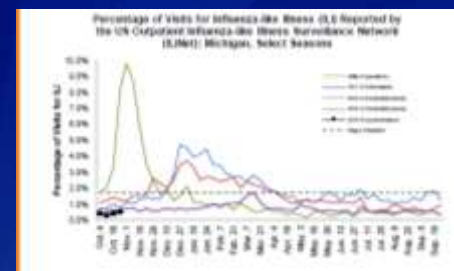
**79,000**



or all the fans in a FULL NFL stadium

get vaccinated

Data source: MMWR, December 13, 2013





## Influenza Vaccine Strains 2014-2015

- ❑ No change from last year's vaccine strains
- ❑ Trivalent vaccine will contain:
  - A/California/7/2009 (H1N1)pdm09-like virus
  - A/Texas/50/2012 (H3N2)-like virus
  - B/Massachusetts/2/2012-like virus
- ❑ Quadrivalent vaccine contains the same three strains as in trivalent vaccine plus:
  - B/Brisbane/60/2008-like virus

MMWR Aug 15, 2014; 63(32):691-7

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## Influenza Vaccines for 2014-15

- ❑ Inactivated (IIV3)
  - Age indications vary by product, formulation and presentation
  - Intramuscular or intradermal injection
  - Trivalent
- ❑ Inactivated (IIV4)
  - Age indications vary by product and presentation
  - Intramuscular injection
  - Quadrivalent
- ❑ Live, attenuated vaccine (LAIV4)
  - Intranasal
  - Quadrivalent

MMWR Aug 15, 2014; 63(32):691-7

## Choice of Influenza Vaccine

- ❑ Where more than one type of vaccine is appropriate and available, ACIP has no other preferential recommendation for use of any influenza vaccine product over another
  - quadrivalent vs trivalent
  - high-dose vs standard dose

MMWR Aug 15, 2014; 63(32):691-7

## Influenza Vaccine Products/Presentations 2014-2015

Name	Age Range	# Antigens	Presentation	Route	Type/Abbrev.
Afluria	5 yrs and older	Trivalent	Pre-Filled Syringe	IM	Inactivated IIV3
	18 yrs thru 64 yrs	Trivalent	Multi-Dose Vial		
Fluarix	3 yrs and older	Trivalent	Pre-Filled Syringe	IM	Inactivated IIV3
		Quadrivalent	Pre-Filled Syringe		
FluBlok	18 yrs and older	Trivalent	Single-Dose Vial	IM	Recombinant RIV3
Flucelvax	18 yrs and older	Trivalent	Pre-Filled Syringe	IM	Cell Culture IIV3
FluLaval	18 yrs and older	Trivalent	Multi-Dose Vial	IM	Inactivated IIV3
		Quadrivalent	Pre-Filled Syringe		
FluMist	2 yrs thru 49 yrs	Quadrivalent	Multi-Dose Vial	IM	Inactivated IIV4
Fluvirin	4 yrs and older	Trivalent	Pre-Filled Syringe	Intranasal	Live Attenuated LAIV4
			Multi-Dose Vial		
Fluzone	6 months and older	Trivalent	Pre-Filled Syringe	IM	Inactivated IIV3
		Quadrivalent	Pre-Filled Syringe		
Fluzone High-Dose	65 yrs and older	Trivalent	Single-Dose Vial	IM	Inactivated IIV4
Fluzone Multi-Dose Vial	65 yrs and older	Trivalent	Multi-Dose Vial	IM	Inactivated IIV3
Fluzone Intradermal	18 yrs thru 64 yrs	Trivalent	Pre-Filled Microinjection System	Intradermal (ID)	Inactivated IIV3

## Influenza Vaccination Schedule

- ❑ Annual single dose for all adults without contraindications or precautions
- ❑ IIV dosage for all adults is 0.5 mL
- ❑ LAIV dosage is 0.2 mL (0.1 mL in each nostril)
  - 2 years – 49 years, healthy, not pregnant

## Vaccinating People with Egg Allergy

**Influenza Vaccination of People with a History of Egg Allergy**

1. People with a history of egg allergy who have experienced only mild allergic reactions to egg-based injectable vaccines should receive any influenza vaccine. Because influenza has the potential for serious complications, influenza vaccine (IIV) should be given for all ages 6 months and older. For people with severe allergic reactions to egg, the following steps should be followed:

2. Influenza vaccine should be administered by a healthcare provider who is familiar with the person's medical history and has the capability to manage severe allergic reactions. The vaccine should be given in a setting where appropriate medical support is available. The vaccine should be given in a setting where appropriate medical support is available. The vaccine should be given in a setting where appropriate medical support is available.

3. People who have had a severe allergic reaction to egg should receive influenza vaccine (IIV) in a setting where appropriate medical support is available. The vaccine should be given in a setting where appropriate medical support is available. The vaccine should be given in a setting where appropriate medical support is available.

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[www.immunize.org/catg.d/p3094.pdf](http://www.immunize.org/catg.d/p3094.pdf)



## Influenza Vaccine and Pregnant Women

- For 2013–14 influenza season, influenza vaccination rate for pregnant women was 52.2%
- Women receiving a clinician recommendation and offer of influenza vaccination had higher vaccination coverage compared with women who reported receiving only a recommendation but no offer or reported receiving no recommendation, *even among those who reported having a negative attitude toward efficacy, safety of influenza vaccination, or no concern about influenza infection*  
MMWR 63(37):816-821

## Vaccinating Mom Also Protects the Baby



## Influenza Vaccination Coverage 2013-14 Healthcare Personnel

Overall – 75.2%

Setting		Occupation	
Hospital	89.6%	Physicians	92.2%
LTC	63.0%	Nurses	90.5%
		NPs & PAs	89.6%
		Pharmacists	85.7%
		Other clinical personnel	87.4%
		Assistants & aides	57.7%
		Nonclinical personnel	68.6%

MMWR, September 19, 2014 / 63(37):805-811

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MMWR, September 19, 2014 / 63(37):805-811

## Influenza Vaccination and HCP

- Coverage was highest among HCP working in settings with flu vaccination requirements (97.8%)
- Coverage by workplace setting was lowest for those working in LTC settings (63.0%)
- Vaccination of HCP in LTC settings is extremely important because:
  - People 65 years and older are at greater risk of serious complications from the flu.
  - Flu vaccine effectiveness is generally lowest in the elderly, making vaccination of close contacts even more critical.
  - Multiple studies have demonstrated health benefits to patients, including reduced flu-related complications and reduced risk of death, with vaccination of HCP in LTC settings.

<http://www.cdc.gov/flu/healthcareworkers.htm>

## CDC Patient Resources



<http://www.cdc.gov/flu/freeresources/print.htm>



## Michigan for Provider and Patients Resources



[www.michigan.gov/immunize](http://www.michigan.gov/immunize)

## Vaccinating People with Egg Allergy

### Influenza Vaccination of People with a History of Egg Allergy



[www.immunize.org/catg.d/p3094.pdf](http://www.immunize.org/catg.d/p3094.pdf)

## ZOSTER VACCINATION



[www.cdc.gov/mmwr/PDF/rr/rr5705.pdf](http://www.cdc.gov/mmwr/PDF/rr/rr5705.pdf)

## ACIP Recommendations for Zoster Vaccine

- ❑ Adults 60 years and older should receive a single dose of zoster vaccine
- ❑ Need for booster dose or doses not known at this time
- ❑ A history of herpes zoster should not influence the decision to vaccinate

MMWR 2008;57(RR-5)

## Zoster Vaccine

- ❑ FDA licensed for adults 50-59 years of age
- ❑ Routine vaccination of adults younger than 60 years NOT recommended by ACIP
- ❑ Rationale
  - Burden of complications highest in persons older than 60 years
  - No recommendation for a booster dose

## Zoster Vaccine

- ❑ It is not necessary to inquire about chickenpox or test for varicella immunity before administering zoster vaccine
- ❑ Persons 60 years of age and older can be assumed to be immune\* regardless of their recollection of chickenpox

MMWR 2008;57(RR-5)

\*For the purpose of establishing eligibility for zoster vaccine



## What Do You Think?



Your 66-year-old patient tells you she had herpes zoster (shingles) 4 years ago. Should you administer zoster vaccine?

- A. Yes – Text IZ3Y to 22333
- B. No – Text IZ3N to 22333

Note: Message and Data Rates May Apply [Poll Results](#)

## HEPATITIS B AND HEALTHCARE PERSONNEL



[www.cdc.gov/mmwr/pdf/rr/r6210.pdf](http://www.cdc.gov/mmwr/pdf/rr/r6210.pdf)

## Hepatitis B Vaccine and Healthcare Personnel (HCP)

- Management of HCP who have written documentation of a complete series of hepatitis B vaccine doses in the past who were not tested for antibody response following the vaccination series and who now test negative for anti-HBs
  - administer 1 dose of hepatitis B vaccine then test for anti-HBs 1 to 2 months later
  - If anti-HBs <10mIU/L, give 2 more doses to complete a second series, then test again 1 to 2 months later.
  - If negative, manage as hepatitis B vaccine non-responder

MMWR 2013;62(RR-10):1-19

## Hepatitis B Vaccine and Healthcare Personnel (HCP)

- Post-exposure prophylaxis
  - when the HBsAg status of the source patient is unknown the exposed, unvaccinated, or incompletely vaccinated HCP should be managed as if the source patient were HBsAg positive
  - include a dose of HBIG in addition to starting or completing the vaccination series for all exposures where the HBsAg status of the source is unknown

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TABLE 3. Postexposure management of health-care personnel after occupational percutaneous and mucous exposure to blood and body fluids, by health-care personnel's hepatitis vaccination and response status

Health-care personnel's vaccination status and response status	Postexposure testing		Postexposure prophylaxis		Post-exposure management
	Time to patient (HBIG)	HBsAg test result	HBIG	Vaccine	
Unvaccinated; exposed after completing series (≥3 doses)	—	—	No action needed	—	—
Unvaccinated; not exposed after 3 doses	Positive/unknown	—	HBIG as reported by 1 month	—	No
	Negative	—	No action needed	—	—
Response unknown after 3 doses	Positive/unknown	—	HBIG as reported by 1 month	—	No
	Negative	—	No action needed	—	—
	Response unknown	—	No action needed	—	—
Unvaccinated; previously vaccinated or vaccine status	Positive/unknown	—	HBIG as reported by 1 month	—	No
	Negative	—	No action needed	—	—

Abbreviations: HBIG = hepatitis B immune globulin; HBsAg = hepatitis B surface antigen; anti-HBs = antibody to hepatitis B surface antigen; HBIG = hepatitis B immune globulin. HBIG should be administered immediately or soon as possible after exposure when indicated. The effectiveness of HBIG when administered > 1 day after administration is unclear. If a health-care personnel is exposed to blood or body fluids and is not vaccinated, the effectiveness of HBIG is uncertain. (Should be performed) 1–2 months after the last dose of the hepatitis vaccine series (and 1–2 months after administration of HBIG to avoid detection of previously administered HBIG using a quantitative restriction algorithm of the protein immunoassay of anti-HBs (anti-HBs)).

A responder is defined as a person with anti-HBs > 10 IU/L (10 mIU/L) after 1–2 doses of hepatitis vaccine.

A non-responder is defined as a person with anti-HBs < 10 IU/L (10 mIU/L) after 1–2 doses of hepatitis vaccine.

HBIG status: anti-HBs < 10 IU/L (10 mIU/L), or who are unvaccinated or incompletely vaccinated, and maintain exposure to a source patient who is HBsAg positive (the unknown HBsAg status, through ongoing testing for HBsAg status as soon as possible after exposure, and follow-up testing approximately 1 month later, to determine the status of the source patient's HBsAg status).

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## ADDITIONAL ADULT IMMUNIZATION RESOURCES





## Resources

- ❑ CDC Vaccines for Adults  
[www.cdc.gov/vaccines/hcp/patient-ed/adults/](http://www.cdc.gov/vaccines/hcp/patient-ed/adults/)
- ❑ CDC Influenza [www.cdc.gov/flu](http://www.cdc.gov/flu)
- ❑ National Foundation for Infectious Diseases (NFID) [www.nfid.org](http://www.nfid.org)
- ❑ ACOG [www.immunizationforwomen.org](http://www.immunizationforwomen.org)

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